IT IS CLAIMED:

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- 1. An isolated polynucleotide capable of encoding a polypeptide having substantial sequence identity to the sequence SEQ ID NO: 2 and characterized by (i) enhanced expression in mammalian central nervous tissue in response to synaptic activation, and (ii) a PDZ-like domain coding region.
 - 2. The isolated polynucleotide of claim 1, wherein said sequence identity is at least about 80%.
- 3. The isolated polynucleotide of claim 1, wherein said polynucleotide has the sequence SEQ 10 ID NO: 1.
 - 4. An isolated polypeptide, characterized by (i) enriched expression during synaptic activity in mammalian brain, (ii) presence of a PDZ-like binding domain, and (iii) a sequence that is at least 80% identical to SEQ ID NO: 2.
 - 5. The isolated polypeptide of claim 4, which further exhibits an ability to selectively bind to a synaptic membrane protein having a C-terminal peptide region selected from the group consisting of SSSL and SSTL.
- The isolated polypeptide of claim 4, wherein said sequence identity is at least about 80%.
 - 7. The isolated polypeptide of claim 6, wherein said polypeptide has the sequence SEQ ID NO: 2.
 - 8. A vector which contains a polynucleotide capable of encoding a polypeptide having at least about 80% sequence identity to the sequence SEQ ID NO: 2 and characterized by enhanced expression in central nervous tissue in response to synaptic activation.
- 30 9. The vector of claim 8, wherein said polynucleotide has the sequence SEQ ID NO: 1.
 - 10. A method of selecting a compound that interferes with binding of a synaptic activation protein to a cellular binding protein in the mammalian central nervous system, comprising
- adding a test compound to a reaction mixture containing (i) an isolated synaptic activation protein having substantial sequence identity to a polypeptide having the sequence SEQ ID NO: 2, (ii) an isolated binding protein to which said synaptic activation protein binds, and (iii) means for detecting

binding between said synaptic activation protein and said binding protein; measuring binding between said synaptic activation protein and said binding protein; and

selecting said compound if the measured binding is greater than or less than binding measured in the absence of said test compound.

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- 11. The method of claim 10, wherein said binding protein is a metabotropic glutamate receptor which includes a sequence selected from the group consisting of SSSL and SSTL.
- The method of claim 11, wherein said mGluR binding protein is expressed in cells, and said binding between said receptor and said binding protein is measured by measuring phosphoinositidase C (PI-PLC) activity in said cells.